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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/762,152	02/01/2001	Chris C. Miller	RR-473PCT/US	1414
7590 02/23/2005			EXAMINER	
SAMUEL N. Tiu, Esq.			HAGHIGHATIAN, MINA	
SIDLEY AUSTIN BROWN & Wood 555 West Fifth Street			ART UNIT	PAPER NUMBER
Los Angeles, CA 90013-1010			1616	

DATE MAILED: 02/23/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

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	Application No.	Applicant(s)
	09/762,152	MILLER, CHRIS C.
Office Action Summary	Examiner	Art Unit
	Mina Haghighatian	1616
The MAILING DATE of this communication a	appears on the cover sheet with	h the correspondence address
Period for Reply A SHORTENED STATUTORY PERIOD FOR REF THE MAILING DATE OF THIS COMMUNICATION - Extensions of time may be available under the provisions of 37 CFR after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a relif NO period for reply is specified above, the maximum statutory perions and the period for reply within the set or extended period for reply will, by state Any reply received by the Office later than three months after the material patent term adjustment. See 37 CFR 1.704(b).	N. 1.136(a). In no event, however, may a reprepay within the statutory minimum of thirty od will apply and will expire SIX (6) MONT tute, cause the application to become ABA	ply be timely filed (30) days will be considered timely. HS from the mailing date of this communication. NDONED (35 U.S.C. § 133).
Status		
1) Responsive to communication(s) filed on 21	October 2004.	
2a) This action is FINAL . 2b) ⊠ T	his action is non-final.	•
3) Since this application is in condition for allow closed in accordance with the practice under		
Disposition of Claims		
4) ☐ Claim(s) 70-99 is/are pending in the applica 4a) Of the above claim(s) is/are withd 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) 70-99 is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) are subject to restriction and	lrawn from consideration.	
Application Papers		
 9) The specification is objected to by the Exam 10) The drawing(s) filed on 01 February 2001 is/Applicant may not request that any objection to the Replacement drawing sheet(s) including the corr 11) The oath or declaration is objected to by the 	/are: a)⊠ accepted or b)□ o he drawing(s) be held in abeyand rection is required if the drawing(s	ce. See 37 CFR 1.85(a). s) is objected to. See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority documed 2. Certified copies of the priority documed 3. Copies of the certified copies of the papplication from the International Buret * See the attached detailed Office action for a light series.	ents have been received. ents have been received in Ap riority documents have been r eau (PCT Rule 17.2(a)).	oplication No received in this National Stage
AM-21-2-24/2)		
Attachment(s) 1) Notice of References Cited (PTO-892)	4) TI Intention St	ummary (PTO-413)
 Notice of References Cited (PTO-032) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/Paper No(s)/Mail Date 	Paper No(s)	//Mail Date formal Patent Application (PTO-152)

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DETAILED ACTION

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 10/21/04 has been entered.

Applicant's submission of amendments, remarks, Declaration of Dr. Neil Macintyre under 37 C.F.R. 1.132, and extension of time filed on 10/21/04 are also acknowledged.

Claim Rejections - 35 USC § 102

The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

f) he did not himself invent the subject matter sought to be patented.

Claims 70-99 are rejected under 35 U.S.C. 102(f) because the applicant did not invent the claimed subject matter.

It appears that the instant invention was not invented by Mr. Chris Miller alone. In light of the letter of Mr. Miller addressed to Dr. Long, dated November 22, 1998, and submitted to the United State Patent and Trademark Office, it is concluded that Mr. Miller was not the sole inventor of the instant invention, as claimed.

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(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 70 and 74-77 are rejected under 35 U.S.C. 102(b) as being anticipated by Green et al (WO 9509612).

Green teaches compositions capable of releasing nitric oxide and therapeutic methods of use thereof for the treatment of microorganism-related disease states. It is discloses that **direct delivery** of nitric oxide **gas** <u>kills intracellular pathogens</u> such as *Mycobacterium tuberculosis*. An ability to <u>specifically</u> deliver compounds capable of releasing nitric oxide to the <u>desired site of infection</u> within the macrophage would greatly enhance killing of intracellular pathogens (page 5, lines 6-13).

Green discloses a method of inhibiting the proliferation of parasites, fungi, bacteria and other proliferating cells or organisms (page 7, lines 30-34). Also disclosed is that the nitric oxide releasing compounds, alone or in combination with other suitable components, can be made into aerosol formulations to be administered via inhalation (page 23, lines 7-10). Administration through the aerosol route is highly beneficial to humans or animals with pulmonary infections. Various bacterial, protozoan, fungal, viral and parasitic infections of the respiratory system that involve macrophages are attacked in this fashion (page 29, lines 15-25).

Green discloses that the dose administered to an animal, particularly a human, should be sufficient to effect a therapeutic response in the animal over reasonable time frame (page 24, line 30 to page 24, line 12).

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Claims 70-78 are rejected under 35 U.S.C. 102(b) as being anticipated by Bathe et al (5,558,083).

Bathe et al teaches a nitric oxide delivery system that is useable with any of a variety of gas delivery system that provides <u>breathing gas to a patient</u>. **Nitric oxide** in a <u>diluent gas is delivered from a gas control valve</u> (see abstract). Bathe discloses that the actual administration of NO is generally carried out by its introduction into the patient as a gas along with other normal inhalation gases given to the patient. Such commercially available supplies are provided in cylinders under pressure (col. 1, lines 20-34). The concentrations at or lower than 150 ppm are suggested (col. 1, lines 35-38).

Bathe discloses that nitric oxide is delivered using systems such as <u>ventilation</u>, where the NO is introduced by means of a gas proportioning device that provides a <u>continuous flow</u> to the patient. The invention includes a flow transducer that senses the flow of gas from the gas delivery system (col. 2, lines 13-30).

Claim Rejections - 35 USC § 103

The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

Claims 70-94 and 97 are rejected under 35 U.S.C. 103(a) as being unpatentable over Bathe et al (5,558,083) in view of Green et al (WO 9509612).

Bathe et al and Green et al, were discussed above. Bathe et al teach a method of delivering nitric oxide gas to patients safely and effectively and Green et al teach the

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beneficial effects of killing the microorganisms of the respiratory system by exposing them directly to nitric oxide gas.

It would have been obvious to a person of ordinary skill in the art at the time the invention was made to have combined the flow-controlled and diluted delivery of nitric oxide gas to patients for respiratory disorders with the teachings of Green et al on the effects of nitric oxide on killing and inhibiting the proliferation of extracellular microorganisms within the respiratory system of an animal because as disclosed nitric oxide can be very beneficial at therapeutic and safe concentration levels and where its delivery is monitored and controlled.

Claims 70-99 are rejected under 35 U.S.C. 103(a) as being unpatentable over Zapol et al (5,873,359) in view of Green et al (WO 9509612).

Zapol et al teach a method of treating or preventing bronchoconstriction or reversible pulmonary vasoconstriction in a mammal, which method includes causing the mammal to inhale a therapeutically-effective concentration of gaseous nitric oxide or a therapeutically-effective amount of a nitric oxide releasing compound, and an inhaler device containing nitric oxide gas and/or nitric oxide-releasing compound (see abstract and summary of the invention).

Zapol discloses that the concentration of nitric oxide is at least 1 ppm, and more preferably at least 20 ppm and most preferably at least 80 ppm (see col. 4, lines 50-59). It is also disclosed that the Occupational Safety and Health Administration (OSHA) has

set the time-weighted average inhalation limit for NO at 25 ppm for 10 hours (col. 3, lines 50-55), and that NO may be administered to mammals at a concentration of from 1 ppm to 40 ppm in air, pure oxygen or another suitable gas or gas mixture, for as along as needed (col. 11, lines 55-61). Zapol discloses that the nitric oxide is inhaled as a mixture including nitric oxide, oxygen and nitrogen gases (col. 4, lines 60-64). Zaplo lacks disclosure on the effects of nitric oxide on microorganisms in the respiratory tract.

Green et al, discussed above, teach the beneficial effects of killing the microorganisms of the respiratory system by exposing them to nitric oxide gas.

It would have been obvious to a person of ordinary skill in the art at the time the invention was made to have combined the method of safely and effectively delivering diluted nitric oxide gas to patients for respiratory disorders with the teachings of Green et al on the effects of nitric oxide on killing and inhibiting the proliferation of extracellular microorganisms within the respiratory system of an animal because as disclosed nitric oxide can be very beneficial at therapeutic and safe concentration levels and where its delivery is monitored and controlled.

Response to Arguments

Applicant's arguments with respect to claims 70-84 have been considered but are most in view of the new ground(s) of rejection. However, since Green et al is still one of

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the references in the new rejection, arguments regarding the said reference will be addressed.

Applicant argues that Green does not disclose the inhalation of nitric oxide gas for killing, inhibiting or suppressing pathogenic microorganisms, but teaches use of generating compounds. Applicant also believes that Green is teaching against the use of nitric oxide gas. This is not commensurate with the scope of the claims. The instant claims do not require a delivery of nitric oxide gas directly to the respiratory system. In fact claims 70 and 79 recite "source of nitric oxide" and claim 76 recites "a nitric oxide substrate source containing a compound capable of producing nitric oxide". The specification also discloses that "nitric oxide source may be a compound, composition or substance producing nitric oxide" (see page 6, lines 7-8).

Applicant argues that Green does not teach the delivery of gaseous nitric oxide through inhalation. This is not commensurate with the scope of the claims. Instant claims require a source of nitric oxide and Green is disclosing a compound releasing nitric oxide gas directly onto the microorganism, which clearly meets the limitation of instant claims. Applicant insists that the instant claims are "limited to the inhalation of nitric oxide gas". This is not correct. Instant claims, especially claim 70 requires "Delivering the nitric oxide gas to the animal's respiratory tract THROUGH inhalation". In response it is disclosed that Green also has inhalation of compounds which <u>Deliver</u> nitric oxide gas to the animal's respiratory tract. Claim 70 also requires that "the inhalation of nitric oxide gas RESULTS in direct exposure of nitric oxide gas to the

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microorganism". Green also discloses that the generating compounds release nitric oxide gas directly onto the microorganisms.

Dr. MacIntyre,s declaration has also been fully considered. It is considered persuasive with regard to claims 79-84, but not persuasive with regard to claims 70-78. Dr. MacIntyre describes Green et al's reference and compares green's disclosure with the instant invention. However Dr. MacIntyre is not considering the scope of the claims. The Office however, is examining the instant CLAIMS and Green is still considered a relevant art for claims 70 and 74-77 under anticipation and claims 70-94 and 97 under obviousness.

However, in response to the Applicant's remarks and the Declaration, it appeared that the rejection should be altered to constitute Bathe et al as the primary reference and Green et al as the supporting prior art.

Bathe et al is clearly teaching a method of safely and effectively administering diluted nitric oxide gas directly and via inhalation to a patient's respiratory system.

Green is relied upon its teaching on the beneficial effects of nitric oxide for killing and inhibiting proliferation of microorganisms within the respiratory system.

It is considered that a combination of Bathe et al and Green et al provides sufficient disclosure to one of ordinary skill in the art to make and use the invention as claimed.

For further clarity it is also disclosed that although Bathe et al does not specifically disclose a method of killing or inhibiting the proliferation of extracellular microorganisms within the respiratory tract of an animal, it is considered that the

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disclosure of Bathe et al on administration of nitric oxide gas to a patient's respiratory system meets such limitations of the claims. This is so because all patients are considered to have a number of microorganisms in their respiratory system, and while the scope of the claims 70-78 does not specifically limit the method to patients with infections, it is considered that Bathe et al is inherently meeting the limitations.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Mina Haghighatian whose telephone number is 571-272-0615. The examiner can normally be reached on core office hours.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary L. Kunz can be reached on 571-272-0887. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

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Mina Haghighatian February 09, 2005 SUPERVISORY PATENT EXAMINER